AMENDMENTS TO THE CLAIMS:

Please amend claims 28, 37, 38, and 40 as shown on the following pages. Material inserted is indicated by underlining (insertion) and material deleted is indicated by strike-out (~~deletion~~).

1-24. (Cancelled)

25. (Previously Presented) A stable pharmaceutical composition comprising a mixture of

(i) an ibuprofen medicament;

(ii) 0.1 to 20% by weight of a domperidone medicament based on the total weight of the composition; and

(iii) a carrier material

characterized in that the carrier material is substantially free of povidone and comprises at least one diluent combined with at least one disintegrating agent excluding

(a) a compressed tablet comprising granulated ibuprofen and a carrier - material consisting essentially of either maize starch at 35-38% total tablet weight in combination with dried maize starch at 3-4% total tablet weight or microcrystalline cellulose at 10-11% total tablet weight in combination with croscarmellose sodium at 14-16% total tablet weight and pre-gelled starch at 10% total tablet weight;

(b) a direct compression tablet comprising a carrier material consisting essentially of microcrystalline cellulose at 8-11% total tablet weight and lactose at 5-6% total tablet weight;

(c) a hard gelatin capsule comprising a carrier consisting essentially of maize starch at 15-20% total capsule contents weight in combination with pre-gelled starch at 5-6% total capsule contents weight.

26. (Previously Presented) A composition according to claim 25 characterised by comprising a granulating agent present to an extent of up to 10% of total tablet weight.

27. (Previously Presented) A composition according to claim 25 comprising a granulating agent consisting essentially of one or more of the following:

a polymeric granulating agents selected from natural gums, synthetic gums and cellulose materials; a sugar granulating agent; a starch granulating agent.

28. (Currently Amended) A composition according to claim ~~38~~ 27 characterised in that the granulating agent is a cellulose derivative ~~hydroxypropyl cellulose or hydroxypropyl methylcellulose~~.

29. (Previously Presented) A composition as claimed in claim 25 in the form of a directly compressed tablet composition comprising:

(i) an ibuprofen medicament;

(ii) a domperidone medicament; and

(iii) a carrier material,

characterised in that the carrier material is substantially free of povidone and comprises at least one diluent combined with at least one disintegrating agent and a lubricating agent.

30. (Previously Presented). A composition according to claim 25 comprising 20-60% by weight carrier material.

31. (Previously Presented) A composition according to claim 25 wherein the carrier material consists essentially of one or more of the following diluents: microcrystalline cellulose, tricalcium phosphate and lactose.

32. (Previously Presented) A composition according to claim 25 comprising one or more discrete disintegrants.

33. (Previously Presented) A composition according to claim 25 in which the ibuprofen medicament is racemic ibuprofen or S(+)-ibuprofen or the sodium or lysine salts thereof, present to an extent of 50-65% by weight of the composition and the domperidone medicament is domperidone or the maleate salt thereof, present to an extent of 1-5% of the composition.

34. (Previously Presented) A process to prepare a composition according to claim 25 comprising (a) granulating said ibuprofen medicament, optionally with said domperidone medicament, with at least a first portion of said carrier material and a granulating fluid; (b) drying said granules; (c) blending with a lubricating agent and optionally a flow aid to form a homogeneous mixture, and (d) compressing into tablets.

35. (Previously Presented) A process according to claim 34 further comprising a cellulose material as a granulating agent.

36. (Previously Presented) A method of treating migraine which comprises the administration to a patient in need thereof a stable pharmaceutical composition according to claim 25.

37. (Currently Amended) A composition as claimed in claim 25 in the form of a compressed tablet wherein the carrier material comprising a compressed mixture of

(a) a granular component comprising said ibuprofen medicament and at least a first portion of said carrier material; and

(b) a powder component comprising a lubricant material and an optional further portion of said carrier material

said domperidone medicament being present in either of components (a) and (b), characterised in that the carrier material is substantially free of povidone and comprises at least one diluent combined with at least one disintegrating agent.

38. (Currently Amended) A composition according to claim ~~27~~ 28 characterised in that the granulating agent is ~~a cellulose derivative~~ hydroxypropyl cellulose or hydroxypropyl methylcellulose.

39. (Previously Presented). A composition according to claim 30, further comprising up to 15% by weight of a discrete disintegrant material.

40. (Currently Amended) A composition according to claim 32 wherein the discrete disintegrant is selected from the group consisting of ~~comprising~~ croscarmellose sodium and sodium starch glycolate.